Human Skin Uptake of Trichloroethylene from Short-Term Aqueous Exposure

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Accelerator mass spectrometry (AMS), a very sensitive analytical method for measuring radioisotopes, was used to measure *in vitro* dermal uptake of 14 C-trichloroethylene (TCE) at very low aqueous concentrations for exposures lasting from 1 to 60 min. Full-thickness human skin (n=3) used within 48 hr of surgery was exposed to $5.3 \pm 1.8 \,\mu\text{g/L}$ of TCE at 22°C in a two-chamber diffusion cell for 1, 5, 15, 30, and 60 min. 14 C content of tissue cores from the exposed skin was determined by AMS to the femtogram (10^{-15}) level. The data from three individuals were expressed as an area- and concentration-normalized flux, R(t), at time t (h), where R(t) has units of cm (i.e., mL of aqueous solution cleared by net 14 C uptake into each dermal plug analyzed, per cm² of exposed plug surface area). When plotted as a function of timet, the data were somewhat nonlinear (p=0.05). An exponential/saturation model fit was consistent with the data (p = 0.84), and yielded an estimated slope (t) of 0.21 \pm 0.045 cm/h.

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